

De-IDTM Software and caBIG

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Features of De-ID

- Accurate, reliable automated de-identification
- Very simple to install and use
 - Network-level operability; batch processing
- Works with word processing files, tab- or comma-delimited or XML formatted data
- Meets all HIPAA 'safe harbor' guidelines;
 option of limited data sets and custom fields

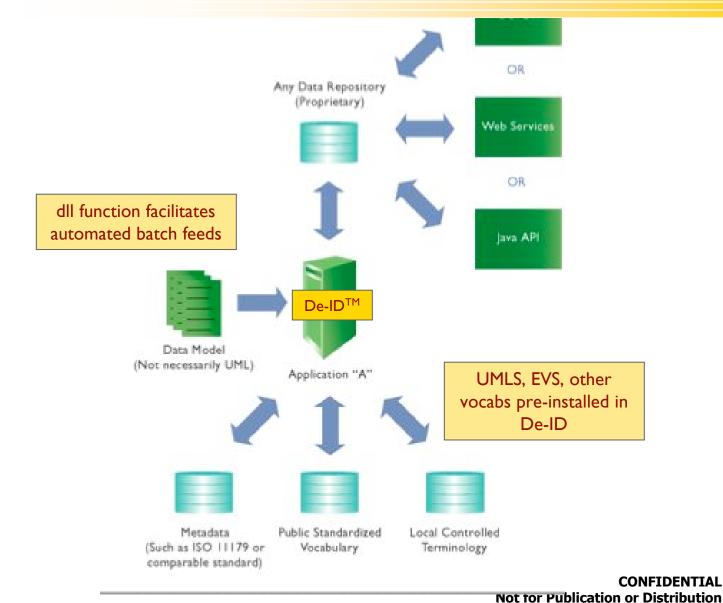


Why Use De-ID™?

Interface Integration	De-ID can be pre-set and 'called' from any application for automated batch processing; can work 'behind the scenes'
Vocabularies/ Terminologies	UMLS, EVS pre-loaded; institution or geography specific glossaries easily loaded
Data elements	Files must be de-identified before leaving 'home' for processing
Information Models	De-identification required before topic mapping or other forms of information modeling

De-ID DATA CORP

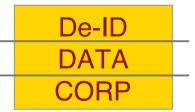
Compatibility Compliance





Compatibility Compliance

- Pre-loaded with UMLS, EVS
 - Can also incorporate specialized acronyms and institutionspecific terminology; identifiers can be managed through customized rule sets
- Automated .dll 'call' from any application for batch file loading
 - Admin-level settings; C/C++header files are created in the subfolder of the install directory.
 - Creates deidentification threads that can be viewed 'in progress' or operate behind the scenes
- Output as independent XML; other file formats as needed
- Outputs automatically fed to content repository or 'next step' application (tagging, parsing)



Benefits of De-ID

- To increase speed and lower costs of "data deposits"
- To maintain consistency of de-identification schema
- To enhance the data value of de-identified records
- Enhances power and usability of traditional de-identification via proxies and offsets for safe-harbor elements, which can be used as data points



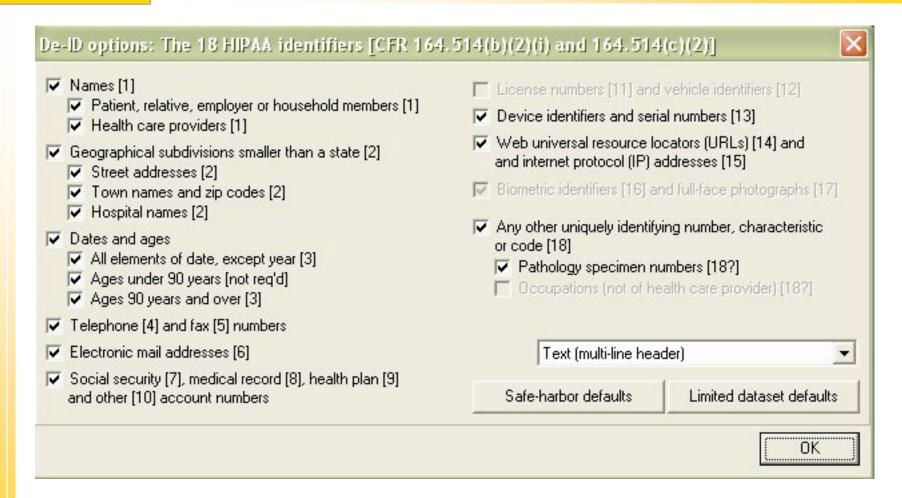
De-ID: A Look Under the Hood

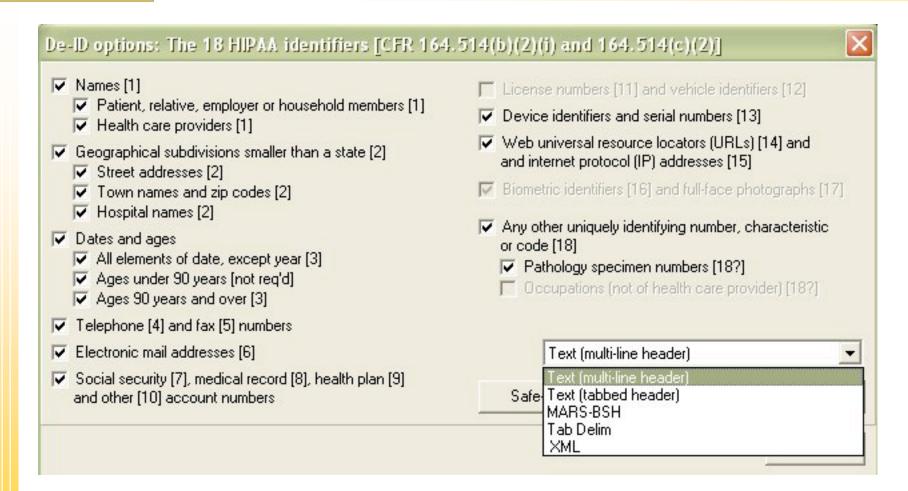
The Manual View

CC 🖭 De-ID / Software for de-identifying narrative medical documents Options Help File Available Header Fields: De-identification/output options Ctrl+D Header Field Type File Type: N/A ▼ Output file (de-identified) ✓ Linkage file 26 Selected Fields: Passphrase: Header Field Type ▼ De-identification log Safe-harbor compliant Text (multi-line header) Begin de-identification De-ID copyright (c) 1999-2004, University of Pittsburgh. All rights reserved. Ver 6.00 (Jan 11 2005)

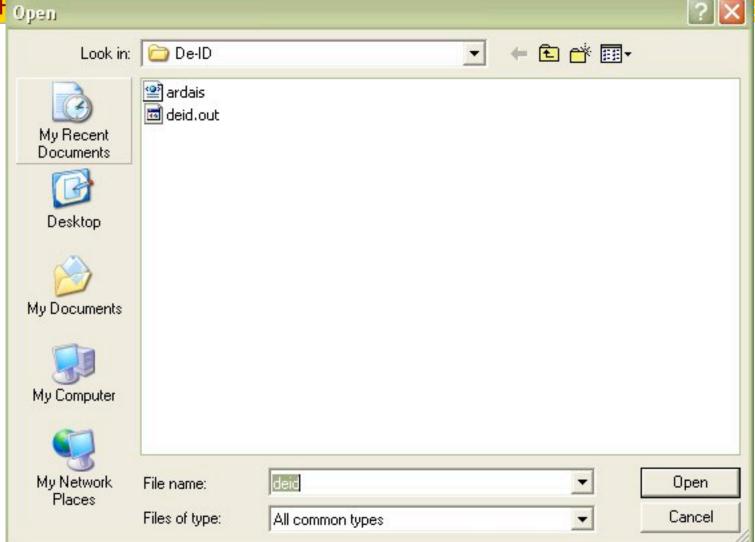
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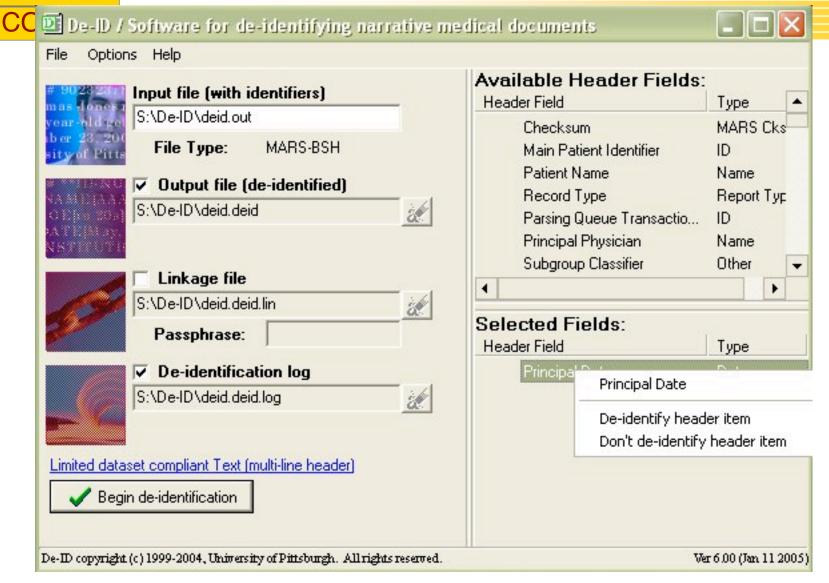


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CC De-ID / Software for de-identifying narrative medical documents File Options Help Available Header Fields: Input file (with identifiers) Header Field Type S:\De-ID\ardais.xml Main Patient Identifier ID File Type: **XML** Record Type Report Type Edit Date Date Output file (de-identified) S:\De-ID\ardais.deid Linkage file 36 S:\De-ID\ardais.deid.lin Selected Fields: Passphrase: Header Field Type De-identification log S:\De-ID\ardais.deid.log Limited dataset compliant Tab Delim Begin de-identification De-ID copyright (c) 1999-2004, University of Pittsburgh. All rights reserved. Ver 6.00 (Jan 11 2005)



De-ID DATA CORP



Text Output 1

```
5 O H
Counters
           Report Type
1,pJYQ49wqYF+h
E O H
[Report de-identified (Safe-harbor compliant) by De-ID v.6.00]
PATIENT HISTORY:
The patient is a " AGE[in 20s] -year-old female with mole changing color and
growing larger.
OSS, *** PATH-NUMBER[1], ** DATE[ May 18 2003]
PRE-OP DIAGNOSIS:
                        Mole of back.
POST-OP DIAGNOSIS: Not given.
/長又現
FINAL DIAGNOSIS:
SKIN, BACK, BIOPSY (OSS, *** PATH-NUMBER[1], ** DATE[ May 18 2003] )
            NON-ULCERATED SUPERFICIAL SPREADING MALIGNANT MELANOMA ARISING IN
AU
A BACKGROUND OF A DYSPLASTIC NEVUS.
       CLARKE'S LEVEL III; 1.7MM 1-2 MITOSIS PER 10 HIGH-POWER FIELD.
Cu
       BRISK MULTIFOCAL TUMOR INFILTRATING LYMPHOCYTES.
       PATHOLOGIC TNM STAGING : DT NM.
**INITIALS
     Pathologist: ** NAME[ WWW ** NAME[ M. M. XXX] , M.D.
     Fellow/Chief Resident: " NAME SSS M. RRR], M.D.
     ** Report Electronically Signed Out **
     By Pathologist: ** NAME[ WWW ** NAME[ M. M. XXX] , M.D.
     ** DATE May 30 2003] 11:56
My signature is attestation that I have personally reviewed the submitted
material(s) and the final diagnosis reflects that evaluation.
```

Text Output 2

ΓIAL

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```
OUTSIDE ACCESSION #:
2 CONSULT SLIDES LABELED *** PATH-NUMBER 1
2 CONSULT BLOCKS LABELED *** PATH-NUMBER 1
OUTSIDE REPORT RECEIVED: Y
CONSULT MATERIAL DESCRIPTION:
Received for consultation from **NAME[M. ZZZ], M.D., are two (2) consult slides
and two (2) consult blocks labeled ***PATH-NUMBER[ 1] from **PLACE,
Pathology Department, "*PLACE, PA, along with an accompanying pathology
JERREE.
/SYR
MICROSCOPIC:
     SYNOPTIC DATA - PRIMARY CUTANEOUS MELANOMA
     Family history of melanoma is unknown.
     Family history of dysplastic nevi is unknown.
TUMOR LOCATION:
                     Back
TYPE OF PROCEDURE:
                        Excisional biopsy
                   Maximum surface diameter of neoplasm: 8 mm
SIZE OF TUMOR:
GROSS ULCERATION:
GROSS SATELLITES:
                      No
HISTOLOGIC TYPE:
                      Superficial spreading
SURFACE ULCERATION:
                     Not present
CLARK'S LEVEL: III
BRESLOW'S THICKNESS:
                          107 mm
ANGIOLYMPHATIC INVASION:
                              None
PERINEURAL INVASION:
                          No
EVIDENCE OF REGRESSION:
MICROSCOPIC SATELLITES:
PREEXISTING NEVUS:
                        Preexisting nevus present
TUMOR INFILTRATING LYMPHOID INFILTRATE:
                                         Marked (Brisk)
MITOTIC RATE:
                  1 / 10 HPF
VASCULARITY:
                  Moderately increased
SURGICAL MARGIN INVOLVEMENT:
                                 Deep margin is free of tumor, Lateral margin
is free of tumor
SENTINEL LYMPH NODE MAPPING:
                                  No
COMPLETION DISSECTION:
T STAGE, PATHOLOGIC:
                          pT2a
N STAGE, PATHOLOGIC:
                          BMX
M STAGE, PATHOLOGIC:
```

Study ID: pJYQ49wgYF+h

Report Type: SP

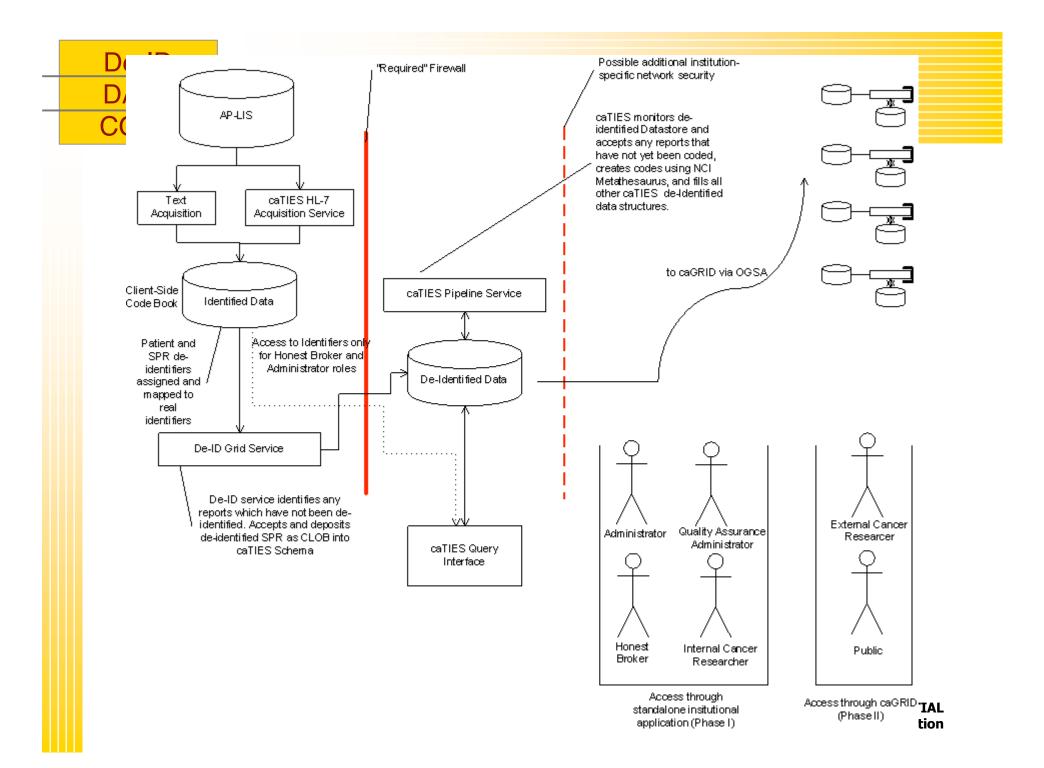
[Report de-identified (Safe-harbor compliant) by De-ID v.6.00] PATIENT HISTORY: The patient is a **AGE[in 20s]-year-old female with mole changing color and growing larger. OSS, ***PATH-NUMBER[1], **DATE[May 18 2003], PRE-OP DIAGNOSIS: Mole of back. POST-OP DIAGNOSIS: Not given. /evm

FINAL DIAGNOSIS:

SKIN, BACK, BIOPSY (OSS, ***PATH-NUMBER[1], **DATE[May 18 2003]) A. NON-ULCERATED SUPERFICIAL SPREADING MALIGNANT MELANOMA ARISING IN A BACKGROUND OF A DYSPLASTIC NEVUS. B. CLARKE'S LEVEL III; 1.7MM 1-2 MITOSIS PER 10 HIGH-POWER FIELD. C. BRISK MULTIFOCAL TUMOR INFILTRATING LYMPHOCYTES. D. PATHOLOGIC TNM STAGING: pT NM. **INITIALS Pathologist: **NAME[WWW **NAME[M. M. XXX], M.D. Fellow/Chief Resident: **NAME[SSS M. RRR], M.D. ** Report Electronically Signed Out ** By Pathologist: **NAME[WWW **NAME[M. M. XXX], M.D. **DATE[May 30 2003] 11:56 My signature is attestation that I have personally reviewed the submitted material(s) and the final diagnosis reflects that evaluation.

OUTSIDE ACCESSION #:

2 CONSULT SLIDES LABELED ***PATH-NUMBER[1] 2 CONSULT BLOCKS LABELED ***PATH-NUMBER[1] OUTSIDE REPORT RECEIVED: Y CONSULT MATERIAL DESCRIPTION: Received for consultation from **NAME[M. ZZZ], M.D., are two (2) consult slides and two (2) consult blocks labeled ***PATH-NUMBER[1] from **PLACE, Pathology Department, **[PLACE] PA, along with an accompanying pathology report. /evm MICROSCOPIC: SYNOPTIC DATA - PRIMARY CUTANEOUS MELANOMA Family history of melanoma is unknown. Family history of dysplastic nevi is unknown. TUMOR LOCATION: Back TYPE OF PROCEDURE: Excisional biopsy SIZE OF TUMOR: Maximum surface diameter of neoplasm: 8 mm GROSS ULCERATION: No GROSS SATELLITES: No HISTOLOGIC TYPE: Superficial spreading SURFACE ULCERATION: Not present CLARK's LEVEL: III BRESLOW'S THICKNESS: 1.7 mm ANGIOLYMPHATIC INVASION: None PERINEURAL INVASION: No EVIDENCE OF REGRESSION: No MICROSCOPIC SATELLITES: No PREEXISTING NEVUS: Preexisting nevus present TUMOR INFILTRATING LYMPHOID INFILTRATE: Marked (Brisk) MITOTIC RATE: 1 / 10 HPF VASCULARITY: Moderately increased SURGICAL MARGIN INVOLVEMENT: Deep margin is free of tumor, Lateral margin is free of tumor SENTINEL LYMPH NODE MAPPING: No COMPLETION DISSECTION: No T STAGE, PATHOLOGIC: pT2a N STAGE, PATHOLOGIC: pNX M STAGE, PATHOLOGIC: pMX -----



Summary

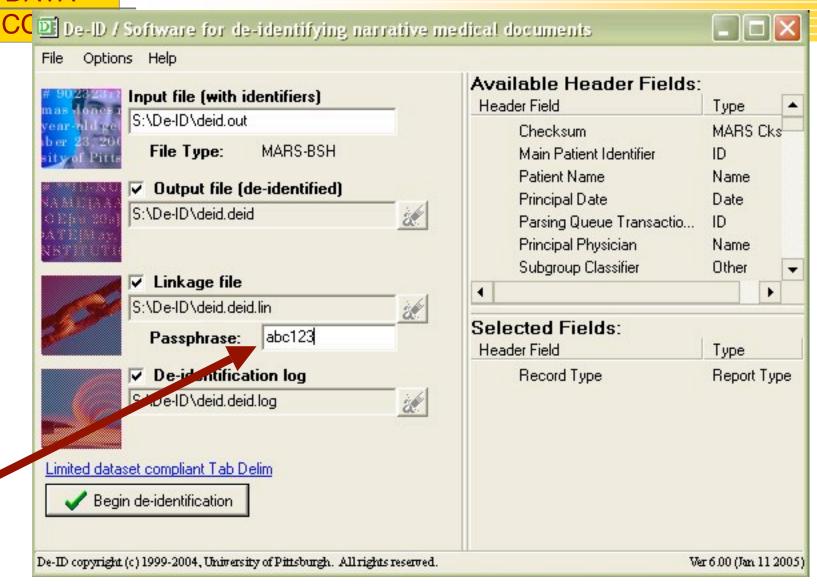
- Network-level operability
- Automated batch feed, output managment
- High volume
- Rapid, reliable
- Leaves original file intact
- Outputs ready for parsing, tagging

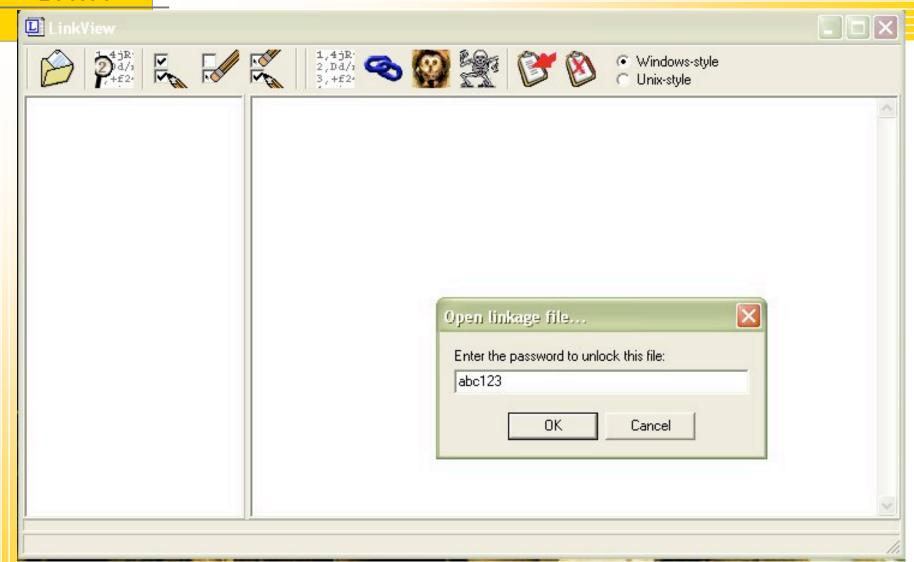


Optional Re-Identification

Re-identification

- Option to create linkage file
- Linkage file independent of de-identified data
- Linkage files are encrypted and password protected





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